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- lymphoblastic leukemia, null cell acute lymphoblastic leukemia, or B cell chronic lymphatic leukemia;
- d) reacts weakly with the human T cell line HJD-1 but does not react with CEM, Laz 191, or HM1;
- e) does not react with the Epstein-Barr virus-transformed human B cell lines
 Laz 007, Laz 156, Laz 256, or SB; and
- f) fixes complement.
- 17. (amended) A method for preparing monoclonal antibody which:
 - a) reacts with essentially all normal human peripheral T cells and cutaneous T lymphoma cells, but not with normal human peripheral B cells, null cells or macrophages;
 - b) reacts with from about 5% to about 10% of normal human thymocytes;
 - c) reacts with leukemic dells from humans with T cell chronic lymphoblastic leukemia but does not react with leukemic cells from humans with T cell acute lymphoblastic leukemia, null call acute lymphoblastic leukemia, or B cell chronic lymphatic leukemia;
 - d) reacts weakly with the human T cell line HJD-1 but does not react with CEM, Laz 191, or HMA;
 - e) does not react with the Epstein-Barr virus-transformed human B cell lines
 Laz 00, Laz 156, Laz 256, or SB; and

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f) fixes complement,

which comprises the steps of:

- i) immunizing mice with E rosette positive purified human/T cells;
- ii) removing the spleens from said mice and making a suspension of the spleen cells;
- iii) fusing said spleen dells with mouse
 myeloma cells in the presence of a
 fusion promoter;
- iv) diluting and culturing the fused
 cells in separate wells in a medium
 which will not support the unfused
 myeloma cells;
- v) evaluating the supernatant in each
 well containing a hybridoma for the
 presence of the desired antibody;
- vi) selecting and cloning hybridomas producing the desired antibody;
- [vii) recovering the antibody from the supernatant above said clones;]
- viii) transferring said clones intraperitoneally into mice; and
 - ix) harvesting the malignant ascites or serum from said mice[.], which ascites or serum contains the desired antibody.

Kindly cancel Claims 4-14 and 19-21.

Add the following new claims: .

Mouse complement-fixing monoclonal antibody which reacts with essentially all normal human peripheral T cells but not with normal human peripheral B cells, null cells, or macrophages.

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A me..... of preparing complement-fixing monoclonal antibody which reacts with essentially all normal human peripheral T cells but not with normal human peripheral B cells, null cells, or macrophages, which comprises culturing the hybridoma ATCC CRL 8001 in a suitable medium and recovering the antibody from the supernatant above said hybridoma.

antibody which reacts with essentially all normal human peripheral. T cells but not with normal human peripheral B cells, null cells, or macrophages, which comprises injecting into a mouse the hybridoma ATCC CRL 8001 and recovering the antibody from the malignant ascites or serum of said mouse.

antibody which reacts with essentially all normal human peripheral T cells but not with normal human peripheral B cells, null cells, ore pared by the method or macrophages, which comprises the steps of:

- i) immunizing mice with E rosette positive purified human T cells;
- ii) removing the spleens from said mice and making a suspension of the spleen cells;
- iii) fusing said spleen cells with mouse
 myeloma cells in the presence of a
 fusion promoter;
- iv) diluting and culturing the fused cells
 in separate wells in a medium which
 will not support the unfused myeloma
 cells;
- v) evaluating the supernatant in each well containing a hybridoma for the presence of antibody to E rosette positive purified T cells;

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- vi) selecting and cloning a hybridoma producing antibody which fixes complement and reacts with essentially all normal human peripheral T cells but not with normal human peripheral B cells, null cells, or macrophages; and
- vii) recovering the antibody from the supernatant above said clones.
- reacts with essentially all normal human peripheral T cells but not with normal human peripheral B cells, null cells, or macroprepared by the method phages, which comprises the steps of:
 - i) immunizing mice with E rosette positive purified human T cells;
 - ii) removing the spleens from said mice and making a suspension of the spleen cells;
 - iii) fusing said spleen cells with mouse
 myeloma cells in the presence of a
 fusion promoter;
 - iv). diluting and culturing the fused cells
 in separate wells in a medium which
 will not support the unfused myeloma
 cells;
 - v) evaluating the supernatant in each well containing a hybridoma for the presence of antibody to E rosette positive purified T cells;
 - vi) selecting and cloning a hybridoma producing antibody which fixes complement and reacts with essentially all normal human peripheral

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